

# CHAPTER 34

## Pulmonary Embolism

### KEY TEACHING POINTS

- In patients with suspected pulmonary embolism the principal role of bedside examination is to identify the patient's overall probability of disease.
- A few *individual* physical findings increase probability of pulmonary embolism—respiratory rate greater than 30/minute, unilateral calf swelling, and parasternal heave—but these findings are infrequent and the increase in probability is only modest.
- By using well-validated scores (e.g., Wells score, revised Geneva score), clinicians can combine risk factors and clinical findings to accurately distinguish patients with low, intermediate, or high probability of pulmonary embolism. This information, combined with quantitative D-dimer measurements, identifies which patients require definitive testing using computed tomography (or ventilation-perfusion lung scanning).

### I. INTRODUCTION

The diagnosis of pulmonary embolism is a difficult one that has frustrated clinicians for more than a century. For example, in up to half of hospitalized patients who die of pulmonary embolism, the diagnosis is not even considered.<sup>1,2</sup> Nowadays, when pulmonary embolism is suspected, the principal role of bedside examination is to determine the patient's overall probability of disease (i.e., low, intermediate, or high probability). This information, in turn, often combined with quantitative D-dimer levels, is used to select which patients should undergo definitive diagnostic testing for thromboembolism by computed tomography (CT) angiography, compression venous ultrasonography, or ventilation-perfusion lung scanning.

### II. THE FINDINGS

Patients with pulmonary embolism present with dyspnea (61% to 83% of patients), pleuritic chest pain (40% to 48% of patients), hemoptysis (5% to 22% of patients), or syncope (4% to 26% of patients).<sup>3-10</sup> Syncope is more common (affecting 20% to 80% of patients) when pulmonary embolism is *massive*, meaning that it obstructs more than half of the pulmonary circulation.<sup>11-13</sup> Ten percent to 35% of patients report a prior history of thromboembolism, and 33% to 42% report calf or thigh pain.<sup>3,5-9</sup>

In recent years, several investigators using multivariate analysis have identified combinations of bedside findings that best identify a patient's overall probability of pulmonary embolism. Two widely studied scores are the **Wells score** (Table 34.1)<sup>14</sup>

TABLE 34.1 Wells Score for Pulmonary Embolism	
Characteristic	Points
<b>RISK FACTORS</b>	
Previous pulmonary embolism or deep venous thrombosis	1.5
Immobilization or surgery in the previous 4 weeks	1.5
Cancer	1
<b>CLINICAL FINDINGS</b>	
Hemoptysis	1
Heart rate > 100/min	1.5
Clinical signs of deep venous thrombosis	3
<b>OTHER</b>	
Alternative diagnosis is less likely than pulmonary embolism	3

Interpretation of total score: 0-1 point, low probability; 2-6 points, moderate probability; 7 or more points, high probability.  
Based upon reference 14.

TABLE 34.2 Revised Geneva Score for Pulmonary Embolism	
Characteristic	Points
<b>RISK FACTORS</b>	
Age >65 years	1
Previous pulmonary embolism or deep venous thrombosis	3
Surgery (under general anesthesia) or fracture (of lower limbs) within 1 month	2
Cancer (active or considered cured <1 year)	2
<b>CLINICAL FINDINGS</b>	
Unilateral leg pain	3
Hemoptysis	2
Heart Rate	
75-94 beats/min	3
≥95 beats/min	5
Pain on palpation of lower-limb deep veins and unilateral edema	4

Interpretation of total score: 0-3 points, low probability; 4-10 points, moderate probability; ≥11 points, high probability.  
Based upon reference 15.

and the revised Geneva score (Table 34.2).<sup>15\*</sup> For each of these scores, the clinician simply adds the points corresponding to each of the independent predictors that are present. The total score determines overall probability, as defined in the footnotes to Tables 34.1 and 34.2. Both scores combine similar risk factors (prior thromboembolism, immobilization, surgery, and cancer) and clinical findings (hemoptysis, tachycardia, and signs of deep venous thrombosis) to arrive at overall clinical

\* The original Geneva score<sup>8</sup> was later revised to remove the patient's arterial blood gas measurement, which is often unavailable.

probability, although the Wells score also considers whether or not an alternative diagnosis is less likely than pulmonary embolism.

### III. CLINICAL SIGNIFICANCE

#### A. INDIVIDUAL FINDINGS

The studies included in [EBM Box 34.1](#) enrolled almost 5000 patients with suspected pulmonary embolism referred to centers having considerable experience with venous thromboembolism. In these studies, only one of five patients suspected of pulmonary embolism actually had the diagnosis.

Very few individual findings help the clinician to distinguish patients with pulmonary embolism from those without it. The only individual symptoms *increasing* the probability of pulmonary embolism are *sudden* dyspnea (likelihood ratio [LR] = 2.4),<sup>6,7</sup> syncope (LR = 2),<sup>4,6</sup> and hemoptysis (LR = 1.9).<sup>3-10†</sup>

The individual physical findings that increase the probability of pulmonary embolism are unilateral calf pain or swelling (LR = 2.5; see [EBM Box 34.1](#)), left parasternal heave (LR = 2.4), respiratory rate of more than 30 breaths/minute (LR = 2), and systolic blood pressure 100 mm Hg or less (LR = 1.9). The presence of wheezes (LR = 0.4) and fever higher than 38°C (LR = 0.5) modestly decrease the probability of pulmonary embolism. The presence or absence of a pulse rate of more than 100/minute as an isolated finding is overall unhelpful (LR = 1.3), although in one study the finding of a pulse less than 90/minute decreased the probability of pulmonary embolism (LR = 0.3).<sup>3</sup>

Other individual findings are unhelpful. Chest wall tenderness is found in 11% to 17% of patients in pulmonary embolism and has a LR that is not significant, emphasizing that this sign is not diagnostic of costochondritis. The presence of hypoxemia, defined either as room air  $pO_2$  less than 80 mm Hg or as increased alveolar-arterial gradient, is also diagnostically unhelpful (both LRs not significant).<sup>3,8,9,30</sup>

#### B. COMBINING FINDINGS TO DETERMINE CLINICAL PROBABILITY OF EMBOLISM

In contrast to the modest accuracy of individual findings, [EBM Box 34.1](#) indicates that a determination of “high probability” by either the Wells score (LR = 7.5) or revised Geneva score (LR = 6.6) markedly increases the probability of pulmonary embolism, whereas a determination of “low probability” by either score decreases it (both LRs = 0.3).

Both scores emphasize that accurate assessment of a patient’s probability combines both risk factors and clinical findings. The probability of embolism is high if the patient has typical signs (e.g., tachycardia, leg swelling) and risk factors (e.g., cancer, immobilization) and lacks an alternative diagnosis. The probability is low if the presentation is atypical, there are no risk factors, and there is a likely alternative diagnosis (e.g., angina, congestive heart failure). Many studies have shown that the probability of pulmonary embolism in patients presenting with both low clinical

† In these studies the following risk factors and symptoms were found just as frequently in patients with embolism as in those without it: female gender, older age, previous heart disease, previous lung disease, estrogen use, recent trauma, dyspnea, chest pain (pleuritic or nonpleuritic), and cough. A few individual risk factors have LRs between 1.3 and 1.9 and thus increase probability a small amount: cancer, recent immobilization, recent surgery, and prior venous thromboembolism.



**EBM BOX 34.1**  
*Pulmonary Embolism\**

Finding (Reference) <sup>†</sup>	Sensitivity (%)	Specificity (%)	Likelihood Ratio <sup>‡</sup> if Finding Is	
			Present	Absent
<b>Individual Findings</b>				
<b>General Description</b>				
Diaphoresis <sup>9</sup>	4	94	NS	NS
Cyanosis <sup>4,9</sup>	1-3	97-100	NS	NS
<b>Vital signs</b>				
Pulse >100/min <sup>6-10,16</sup>	6-43	66-96	1.3	NS
Systolic blood pressure ≤100 mm Hg <sup>8</sup>	8	95	1.9	NS
Temperature >38°C <sup>4,6-9</sup>	1-9	78-98	0.5	NS
Respiratory rate >30/ min <sup>8</sup>	21	90	2.0	0.9
<b>Lung</b>				
Accessory muscle use <sup>4</sup>	17	89	NS	NS
Crackles <sup>3,9,17</sup>	21-59	45-82	NS	NS
Wheezes <sup>6,9,17</sup>	3-31	68-91	0.4	NS
Pleural friction rub <sup>4,9</sup>	1-14	91-99	NS	NS
<b>Heart</b>				
Elevated neck veins <sup>4,9,17</sup>	3-14	92-96	1.7	NS
Left parasternal heave <sup>4,9</sup>	1-5	98-99	2.4	NS
Loud P <sub>2</sub> <sup>3,9</sup>	15-19	84-95	NS	NS
New gallop (S <sub>3</sub> or S <sub>4</sub> ) <sup>3</sup>	30	89	NS	NS
<b>Other</b>				
Chest wall tenderness <sup>4,18</sup>	11-17	79-80	NS	NS
Unilateral calf pain or swelling <sup>5-7,9,10,17,19</sup>	9-52	77-99	2.5	0.8
<b>Combined Findings</b>				
<b>Wells Score<sup>7,20-29</sup></b>				
Low probability, 0-1 points	6-53	30-54	0.3	—
Moderate probability, 2-6 points	38-72	—	1.6	—
High probability, 7 or more points	7-54	90-100	7.5	—

**EBM BOX 34.1***Pulmonary Embolism\*—cont'd*

Finding (Reference) <sup>†</sup>	Sensitivity (%)	Specificity (%)	Likelihood Ratio <sup>‡</sup> if Finding Is	
			Present	Absent
<b>Revised Geneva Score</b> <sup>15,24-27</sup>				
Low probability, 0-3points	1-27	43-85	0.3	—
Moderate probability, 4-10 points	58-69	—	NS	—
High probability, ≥11 points	10-42	96-99	6.6	—

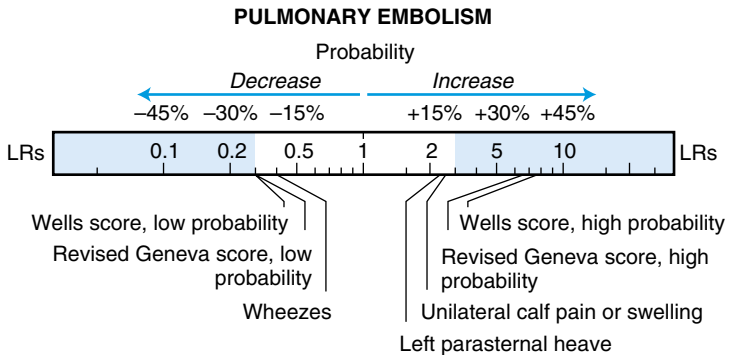
\*Diagnostic standard: For *pulmonary embolism*, pulmonary angiography, CT angiography, or ventilation-perfusion scanning ( $\pm$  compression venous ultrasonography). In eight studies,<sup>15,18,20,21,23,25,27,28</sup> some low-risk patients (i.e., those with negative quantitative D-dimers and low clinical risk) were not tested but instead were followed at least 3 months without anticoagulation; all lacked clinical evidence of thromboembolism.

<sup>†</sup>Definition of findings: for *Wells score* and *revised Geneva score*, see [Tables 34.1 and 34.2](#).

<sup>‡</sup>Likelihood ratio (LR) if finding present = positive LR; LR if finding absent = negative LR.

NS, Not significant.

[Click here to access calculator](#)



probability (using either score) and normal D-dimer levels is so low that further imaging is unnecessary and anticoagulation can safely be withheld.<sup>15,21,23,25,31,32</sup>

The references for this chapter can be found on [www.expertconsult.com](http://www.expertconsult.com).

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## REFERENCES

1. Ryu JH, Olson EJ, Pellikka PA. Clinical recognition of pulmonary embolism: problem of unrecognized and asymptomatic cases. *Mayo Clin Proc.* 1998;73:873–879.
2. Morgenthaler TI, Ryu JH. Clinical characteristics of fatal pulmonary embolism in a referral hospital. *Mayo Clin Proc.* 1995;70:417–424.
3. Hoellerich VL, Wigton RS. Diagnosing pulmonary embolism using clinical findings. *Arch Intern Med.* 1986;146:1699–1704.
4. Hull RD, Raskob GE, Carter CJ, et al. Pulmonary embolism in outpatients with pleuritic chest pain. *Arch Intern Med.* 1988;148:838–844.
5. Kline JA, Nelson RD, Jackson RE, Courtney DM. Criteria for the safe use of D-dimer testing in emergency department patients with suspected pulmonary embolism: a multicenter US study. *Ann Emerg Med.* 2002;39(2):144–152.
6. Miniati M, Monti S, Bottai M. A structured clinical model for predicting the probability of pulmonary embolism. *Am J Med.* 2003;114:173–179.
7. Miniati M, Bottai M, Monti S. Comparison of 3 clinical models for predicting the probability of pulmonary embolism. *Medicine.* 2005;84:107–114.
8. Wicki J, Perneger TV, Junod AF, Bounameaux H, Perrier A. Assessing clinical probability of pulmonary embolism in the emergency ward: a simple score. *Arch Intern Med.* 2001;161:92–97.
9. Stein PD, Beemath A, Matta F, et al. Clinical characteristics of patients with acute pulmonary embolism: data from PIOPEd II. *Am J Med.* 2007;120:871–879.
10. Tsimogianni AM, Rovina N, Porfyridis I, et al. Clinical prediction of pulmonary embolism in respiratory emergencies. *Thromb Res.* 2011;127:411–417.
11. Stein PD, Willis PW, DeMets DL. History and physical examination in acute pulmonary embolism in patients without preexisting cardiac or pulmonary disease. *Am J Cardiol.* 1981;47:218–223.
12. Bell WR, Simon TL, DeMets DL. The clinical features of submassive and massive pulmonary emboli. *Am J Med.* 1977;62:355–360.
13. Sutton GC, Honey M, Gibson RV. Clinical diagnosis of acute massive pulmonary embolism. *Lancet.* 1969;1:271–273.
14. Wells PS, Anderson DR, Rodger M, et al. Derivation of a simple clinical model to categorize patients probability of pulmonary embolism: increasing the models utility with the SimpliRED D-dimer. *Thromb Haemost.* 2000;83:416–420.
15. Le Gal G, Righini M, Roy PM, et al. Prediction of pulmonary embolism in the emergency department: the revised Geneva score. *Ann Intern Med.* 2006;144:165–171.
16. Kline JA, Corredor DM, Hogg MM, Hernandez J, Jones AE. Normalization of vital signs does not reduce the probability of acute pulmonary embolism in symptomatic emergency department patients. *Acad Emerg Med.* 2012;19:11–17.
17. Chen JY, Chao TH, Guo YL, et al. A simplified clinical model to predict pulmonary embolism in patients with acute dyspnea. *Int Heart J.* 2006;47:259–271.
18. Le Gal G, Testuz A, Righini M, Bounameaux H, Perrier A. Reproduction of chest pain by palpation: diagnostic accuracy in suspected pulmonary embolism. *Br Med J.* 2005;330:452–453.
19. Stein PD, Henry JW, Gopalakrishnan D, Relyea B. Asymmetry of calves in the assessment of patients with suspected acute pulmonary embolism. *Chest.* 1995;107:936–939.
20. Chagnon I, Bounameaux H, Aujesky D, et al. Comparison of two clinical prediction rules and implicit assessment among patients with suspected pulmonary embolism. *Am J Med.* 2002;113:269–275.
21. Wells PS, Anderson DR, Rodger M, et al. Excluding pulmonary embolism at the bedside without diagnostic imaging: management of patients with suspected pulmonary embolism presenting to the emergency department by using a simple clinical model and D-dimer. *Ann Intern Med.* 2001;135:98–107.
22. Wolf SJ, McCubbin TR, Feldhaus KM, Faragher JP, Adcock DM. Prospective validation of Wells criteria in the evaluation of patients with suspected pulmonary embolism. *Ann Emerg Med.* 2004;44:503–510.

23. Anderson DR, Kovacs MJ, Dennie C, et al. Use of spiral computed tomography contrast angiography and ultrasonography to exclude the diagnosis of pulmonary embolism in the emergency department. *J Emerg Med.* 2005;29(4):399–404.
24. Calisir C, Yavas US, Ozkan IR, et al. Performance of the Wells and revised Geneva scores for predicting pulmonary embolism. *Eur J Emerg Med.* 2008;16:49–52.
25. Klok FA, Kruisman E, Spaan J, et al. Comparison of the revised Geneva score with the Wells rule for assessing clinical probability of pulmonary embolism. *J Thromb Haemost.* 2008;6:40–44.
26. Wong DD, Ramaseshan G, Mendelson RM. Comparison of the Wells and revised Geneva scores for the diagnosis of pulmonary embolism: an Australian experience. *Intern Med J.* 2011;41:258–262.
27. Penaloza A, Verschuren F, Meyer G, et al. Comparison of the unstructured clinician gestalt, the Wells score, and the revised Geneva score to estimate pretest probability for suspected pulmonary embolism. *Ann Emerg Med.* 2013;62:117–124.
28. Penaloza A, Melot C, Motte S. Comparison of the Wells score with the simplified revised Geneva score for assessing pretest probability of pulmonary embolism. *Thromb Res.* 2011;127:81–84.
29. Yap KS, Kalff V, Turlakow A, Kelly MJ. A prospective reassessment of the utility of the Wells score in identifying pulmonary embolism. *Med J Aust.* 2007;187:333–336.
30. Stein PD, Goldhaber SZ, Henry JW, Miller AC. Arterial blood gas analysis in the assessment of suspected acute pulmonary embolism. *Chest.* 1996;109(1):78–81.
31. Leclercq MGL, Lutisan JG, van Marwijk Kooy M, et al. Ruling out clinical suspected pulmonary embolism by assessment of clinical probability and D-dimer levels: a management study. *Thromb Haemost.* 2003;89:97–103.
32. Kruip MJHA, Leclercq MGL, van der Heul C, Prins MH, Buller HR. Diagnostic strategies for excluding pulmonary embolism in clinical outcome studies: a systematic review. *Ann Intern Med.* 2003;138:941–951.